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TERMINAL (ENTER 1, 2, 3, OR ?):2

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NEWS
NEWS
                 "Ask CAS" for self-help around the clock
                 CA/CAplus records now contain indexing from 1907 to the
NEWS
         SEP 09
                 present
NEWS
         DEC 08
                 INPADOC: Legal Status data reloaded
NEWS
         SEP 29 DISSABS now available on STN
         OCT 10
                 PCTFULL: Two new display fields added
NEWS
     6
         OCT 21 BIOSIS file reloaded and enhanced
NEWS
     7
         OCT 28 BIOSIS file segment of TOXCENTER reloaded and enhanced
NEWS 8
NEWS 9 NOV 24 MSDS-CCOHS file reloaded
NEWS 10 DEC 08
                 CABA reloaded with left truncation
        DEC 08
NEWS 11
                 IMS file names changed
         DEC 09
NEWS 12
                 Experimental property data collected by CAS now available
                 in REGISTRY
NEWS 13
         DEC 09
                 STN Entry Date available for display in REGISTRY and CA/CAplus
NEWS 14
        DEC 17
                 DGENE: Two new display fields added
NEWS 15
         DEC 18
                 BIOTECHNO no longer updated
NEWS 16 DEC 19
                 CROPU no longer updated; subscriber discount no longer
                 available
NEWS 17 DEC 22
                 Additional INPI reactions and pre-1907 documents added to CAS
                 databases
NEWS 18 DEC 22
                 IFIPAT/IFIUDB/IFICDB reloaded with new data and search fields
NEWS 19
        DEC 22
                 ABI-INFORM now available on STN
NEWS 20
        JAN 27
                 Source of Registration (SR) information in REGISTRY updated
                 and searchable
NEWS 21 JAN 27
                 A new search aid, the Company Name Thesaurus, available in
                 CA/CAplus
        FEB 05
NEWS 22
                 German (DE) application and patent publication number format
                 changes
             DECEMBER 28 CURRENT WINDOWS VERSION IS V7.00, CURRENT
NEWS EXPRESS
              MACINTOSH VERSION IS V6.0b(ENG) AND V6.0Jb(JP),
              AND CURRENT DISCOVER FILE IS DATED 23 SEPTEMBER 2003
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              STN Operating Hours Plus Help Desk Availability
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              General Internet Information
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              Welcome Banner and News Items
NEWS PHONE
              Direct Dial and Telecommunication Network Access to STN
NEWS WWW
              CAS World Wide Web Site (general information)
```

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* * * STN Columbus

FILE 'HOME' ENTERED AT 10:07:11 ON 12 FEB 2004

=> file reg

COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 0.21 0.21

FULL ESTIMATED COST

FILE 'REGISTRY' ENTERED AT 10:07:20 ON 12 FEB 2004 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2004 American Chemical Society (ACS)

Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

HIGHEST RN 649538-27-2 STRUCTURE FILE UPDATES: 11 FEB 2004 DICTIONARY FILE UPDATES: 11 FEB 2004 HIGHEST RN 649538-27-2

TSCA INFORMATION NOW CURRENT THROUGH JULY 14, 2003

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at: http://www.cas.org/ONLINE/DBSS/registryss.html

Uploading c:\09847654-4.str

STRUCTURE UPLOADED L1

=> d 11

L1 HAS NO ANSWERS

L1

STR Ak-O OH OH

Structure attributes must be viewed using STN Express query preparation.

=> s l1 sss sam SAMPLE SEARCH INITIATED 10:07:52 FILE 'REGISTRY' SAMPLE SCREEN SEARCH COMPLETED -3274 TO ITERATE

1000 ITERATIONS 30.5% PROCESSED INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED) SEARCH TIME: 00.00.01

21 ANSWERS

FULL FILE PROJECTIONS: ONLINE **COMPLETE** BATCH **COMPLETE** PROJECTED ITERATIONS: PROJECTED ANSWERS:

62049 TO 68911 878 TO 1872

L2

21 SEA SSS SAM L1

=> d scan

21 ANSWERS REGISTRY COPYRIGHT 2004 ACS on STN L2

IN Phosphoric acid, P,P'-1,2-ethanediyl P,P'-bis(2,2,6,6-tetramethyl-4-

piperidinyl) ester, nickel(2+) salt (1:1) (9CI)

MF C20 H42 N2 O8 P2 . Ni

Ni(II)

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):1

L221 ANSWERS REGISTRY COPYRIGHT 2004 ACS on STN

IN D-myo-Inositol, 4,5-bis(dihydrogen phosphate) 1-[(2R,3R)-11-(3',6'dihydroxy-3-oxospiro[isobenzofuran-1(3H),9'-[9H]xanthen]-5-yl)-6-hydroxy-6oxido-11-oxo-2,3-bis[(1-oxohexadecyl)oxy]-5,7-dioxa-10-aza-6-phosphaundec-1-yl hydrogen phospahte] (9CI)

C65 H99 N O29 P4

Absolute stereochemistry.

PAGE 1-A

HO_

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):1

L2 21 ANSWERS REGISTRY COPYRIGHT 2004 ACS on STN

IN Guanosine, 2'-deoxyguanylyl-(3'→5')-2'-deoxycytidylyl(3'→5')-2'-deoxyguanylyl-(3'→5')-2'-deoxyadenylyloxy[(2S,3S)3-methyl-2-[[5-[(1Z)-phenylazo]benzoyl]amino]-1,3propanediyl]oxyphosphinico-(3'→5')-2'-deoxyguanylyl-(3'→5')thymidylyl-(3'→5')-2'-deoxycytidylyl-(3'→5')-2'-deoxy-,
complex with 2'-deoxycytidylyl-(3'→5')-2'-deoxyguanylyl(3'→5')-2'-deoxyadenylyl-(3'→5')-2'-deoxycytidylyl(3'→5')-thymidylyl-(3'→5')-2'-deoxycytidylyl-(3'→5')2'-deoxyguanylyl-(3'→5')-2'-deoxycytidine (1:1) (9CI)

MF C95 H116 N36 O51 P8 . C76 H98 N29 O46 P7

CM 1

Absolute stereochemistry. Double bond geometry as shown.

PAGE 1-A

PAGE 1-C

 \sim NH₂

PAGE 2-B

PAGE 3-A

Absolute stereochemistry.

PAGE 1-A

но<u></u>

PAGE 1-B

$$\begin{array}{c} & & & & & \\ & & & & \\ & & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & &$$

PAGE 2-B

PAGE 3-A

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):1

L2 21 ANSWERS REGISTRY COPYRIGHT 2004 ACS on STN

CM 1

Absolute stereochemistry.

PAGE 1-A

CM 2

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):end

=> d l1 sss full
L1 HAS NO ANSWERS
'SSS FULL ' IS NOT A VALID STRUCTURE FORMAT KEYWORD
Structure Formats
SIA ---- Structure Image, Attributes, and map table if it contains data. (Default)
SIM ---- Structure IMage.

SAT ---- Structure ATtributes and map table if it contains data.

SCT ---- Structure Connection Table and map table if it contains

data.

SDA ---- All Structure DAta (image, attributes, connection table and map table if it contains data).

NOS ---- NO Structure data.

ENTER STRUCTURE FORMAT (SIM), NOS:end

=> s l1 sss full

FULL SEARCH INITIATED 10:08:54 FILE 'REGISTRY' FULL SCREEN SEARCH COMPLETED - 66898 TO ITERATE

100.0% PROCESSED 66898 ITERATIONS 1425 ANSWERS

SEARCH TIME: 00.00.08

1425 SEA SSS FUL L1

=> file caplus

COST IN U.S. DOLLARS

SINCE FILE TOTAL

ENTRY 156.26

SESSION 156.47

FULL ESTIMATED COST

FILE 'CAPLUS' ENTERED AT 10:09:16 ON 12 FEB 2004 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2004 AMERICAN CHEMICAL SOCIETY (ACS)

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FILE COVERS 1907 - 12 Feb 2004 VOL 140 ISS 7 FILE LAST UPDATED: 11 Feb 2004 (20040211/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s 13 and antimicrobial

8116 L3

49629 ANTIMICROBIAL

3378 ANTIMICROBIALS

50747 ANTIMICROBIAL

(ANTIMICROBIAL OR ANTIMICROBIALS)

L410 L3 AND ANTIMICROBIAL

=> dis l4 1-10 bib abs hitstr

1.4 ANSWER 1 OF 10 CAPLUS COPYRIGHT 2004 ACS on STN

AN 2003:31542 CAPLUS

DN 139:239970

An assessment of fructose-1,6-bisphosphate as an antimicrobial ТT and anti-inflammatory agent in sepsis

ΔII Nunes, Fernanda B.; Graziottin, Cassio M.; Alves Filho, Jose Carlos F.; Lunardelli, Adroaldo; Pires, Melissa G. S.; Wachter, Paulo H.; De Oliveira, Jarbas R.

CS Faculdade de Biociencias, Departamento de Ciencias Fisiologicas, Laboratorio de Pesquisa em Biofisica, Pontificia Universidade Catolica do Rio Grande do Sul, Rio Grande do Sul, Brazil

SO Pharmacological Research (2003), 47(1), 35-41

CODEN: PHMREP; ISSN: 1043-6618

PB Elsevier Science Ltd.

DT Journal

LA English

Tissue lesion mechanisms provoked by sepsis include the infectious AB process, inflammation, and cellular energy deficit. We chose to test fructose-1,6-bisphosphate (FBP) because of its possible anti-inflammatory and antimicrobial actions. Wistar rats were used and divided into three exptl. groups: a control group (n = 10), in which a capsule was introduced into the peritoneum of the animals; a septic group (n = 10), in which a capsule containing non-sterile fecal matter was introduced together with Escherichia coli (1.5 + 109 CFU); and a septic group treated with FBP 500 mg/kg (n = 10). The blood cell tests revealed that levels of leukocytes increased significantly in the septic group when compared to both the septic group treated with FBP and the control group. The blood cultures were 100% pos. in both the septic group and the septic group treated with bisphosphorylated sugar. The antibiogram only revealed an inhibitory halo in the case of the antibiotic ampicillin, there was no such indication for FBP. The anti-inflammatory power of FBP remained at 60% for 5 h in the rats that received the carrageenan injection. What is more, the sugar reduced the levels of ionic calcium in relation to the control group. This data proves the validity of using FBP in the treatment of sepsis, possibly due to its anti-inflammatory rather than antimicrobial action.

IT 488-69-7, Fructose 1,6 bisphosphate

RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(assessment of fructose-1,6-bisphosphate as an antimicrobial and anti-inflammatory agent in sepsis)

RN 488-69-7 CAPLUS

CN D-Fructose, 1,6-bis(dihydrogen phosphate) (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RE.CNT 38 THERE ARE 38 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 2 OF 10 CAPLUS COPYRIGHT 2004 ACS on STN

AN 2003:5224 CAPLUS

DN 138:61095

TI Dentifrice compositions containing antimicrobial enzymes

IN Dana, Frederic

PA Fr

SO U.S. Pat. Appl. Publ., 13 pp., Cont.-in-part of U.S. Ser. No. 872,829, abandoned.

CODEN: USXXCO

DT Patent

LA English

FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
ΡI	US 2003003059	A1	20030102	US 2001-4111	20011115
	FR 2822700	A1	20021004	FR 2001-4614	20010403

```
WO 2003043517
                                           WO 2002-US36659 20021114
                            20030530
                       A2
                            20030918
     WO 2003043517
                       A3
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
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             LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
             PL, PT, RO, RU, SC, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT,
             TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
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             FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF,
             CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
                            20010403
PRAI FR 2001-4614
                       Α
     US 2001-872829
                       В2
                            20010601
     US 2001-4111
                            20011115
                       Α
     The present invention relates to oral care compns. which provide a means
AB
     to deliver actives useful in the prevention, treatment and/or management
     of dental and related tissue conditions, including dental caries, dental
     cavities, microbial flora, tartar, periodontal and related gum disease.
     In addition, the present invention may be used in the healthy maintenance of
     teeth and gums of humans and pets. The present compns. are useful to
     whiten teeth and otherwise favorably impact the cosmetic appeal of the
     teeth and gums of a subject or patient. The inclusion of effective amts.
     of colostrum in dental care compns. provides an unexpectedly high efficacy
     of such formulations in inhibiting, reducing or otherwise preventing
     microbial growth, dental caries, plaque, cavities and gum disease,
     including periodontal disease. The use of colostrum with other enzymes,
     e.g., lysozyme, lactoperoxidase, dextranase, mutanase, cellulase,
     amyloglucosidase, papain, bromelin, lactoferrin, etc., represents a
     particularly preferred embodiment for use in the present invention because
     of the unexpected antimicrobial activity exhibited by the enzyme
     combination. For example, chewable dentifrice tablets were prepared containing
     (by weight) bromochlorophene 0.01-1%, enoxolone 0.1-3%, sodium bicarbonate
     1-5%, silica 1-5%, sorbitol 45-60%, xylitol 5-40%, liver powder 1-15%,
     methionine/cysteine 0.1-3%, Coloring 5 0.001-0.1%, papain/bromelin
     0.01-1%, glucose oxidase/lactoperoxidase 0.01-1%,
     amyloglucosidase/invertase 0.01-1%, and lysozyme/lactoferrin 0.01-1%.
IT
     112084-16-9
     RL: COS (Cosmetic use); BIOL (Biological study); USES (Uses)
        (anti-tartar agent; ingestible dentifrice compns. containing colostrum with
        other antimicrobial enzymes)
RN
     112084-16-9 CAPLUS
CN
     1,1,1-Ethanetriol, bis(dihydrogen phosphate) (9CI) (CA INDEX NAME)
       OH
H2O3PO-C-Me
       OPO3H2
L4
     ANSWER 3 OF 10 CAPLUS COPYRIGHT 2004 ACS on STN
AN
     2002:185692 CAPLUS
DN
     136:236873
TТ
     Protonated antimicrobial compounds
IN
    Dale, Roderic M. K.; Gatton, Steven L.; Arrow, Amy; Thompson, Terry
PA
so
    U.S. Pat. Appl. Publ., 28 pp., Cont.-in-part of U.S. Ser. No. 281,858.
     CODEN: USXXCO
DT
     Patent
    English
FAN.CNT 5
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KIND DATE
                                          APPLICATION NO. DATE
     PATENT NO.
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                                          -----
PΙ
     US 2002032164
                     A1
                           20020314
                                         US 2001-847654
                                                           20010503
                                          US 1998-222009
     US 6211349
                      B1
                           20010403
                                                           19981230
                                          US 1999-281858
     US 6627215
                      B1
                           20030930
                                                           19990331
     WO 2002089581
                     A1
                           20021114
                                          WO 2002-US13910 20020503
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            LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
            PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ,
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                          19981230
PRAI US 1998-222009
                     A2
    US 1999-281858
                      A2
                           19990331
    US 2001-847654
                      Α
                           20010503
os
    MARPAT 136:236873
AB
    The present invention provides protonated compds. X-Y-Z (Y = O, P, C; X, Z
     = end blocking groups preventing degradation of the mol. and providing
     stability) having antimicrobial activity and a sanitizing composition
    comprising a protonated compound and a metal salt of a carboxylic acid.
    protonated compds. and compns. provide efficacious antimicrobial
    activity against resistant strains of bacteria and opportunistic fungi.
    For example, the s.c. administration of compds. Nu-2, Nu-3, Nu-4, and Nu-5
     (12 mg/mL) were effective in attenuating the incidence of infection of
    burn wounds in a mice model, a ribose derivative Nu-4 being the most
    efficacious providing 100% survival.
```

IT 403717-08-8

RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(protonated antimicrobial compds. and compns.)

RN 403717-08-8 CAPLUS

CN Phosphoric acid, P,P'-1,4-butanediyl P,P'-dibutyl ester (9CI) (CA INDEX NAME)

```
L4
     ANSWER 4 OF 10 CAPLUS COPYRIGHT 2004 ACS on STN
AN
     2002:71818 CAPLUS
DN
     136:107271
TI
     Cosmetic emulsions containing hemoglobin and myoglobin as oxygen carriers
     for the natural regeneration of skin in case of oxygen deficiency
     Barnikol, Wolfgang
IN
PA
     SanguiBioTech A.-G., Germany
SO
     PCT Int. Appl., 35 pp.
     CODEN: PIXXD2
DT
     Patent
LΑ
     German
FAN.CNT 1
     PATENT NO.
                     KIND DATE
                                         APPLICATION NO. DATE
                                          -----
ΡI
     WO 2002005754
                    A2
                           20020124
                                         WO 2001-EP7495
                                                          20010629
     WO 2002005754
                    A3
                           20020718
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W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,

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             SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN,
             YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
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             BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                            20020207
                                          DE 2000-10034970 20000719
     DE 10034970
                       A1
     EP 1301169
                                          EP 2001-953186
                       A2
                            20030416
                                                            20010629
             AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
                                           US 2003-333167
     US 2004022839
                            20040205
                                                            20030522
                       A1
PRAI DE 2000-10034970 A
                            20000719
     WO 2001-EP7495
                       W
                            20010629
     The invention relates to an emulsion in the form of a cream or a lotion
AB
     and to the usage as cosmetics. The emulsion has a caring effect on the
     skin and improves the diffusive oxygen supply of the epidermis for the
     purpose of regenerating it and of remedying an oxygen deficiency. The
     oxygen carrier is a native or modified Hb or a Hb/myoglobin mixture The use
     of one or more oily components together with one or more O/W emulsifiers
     improves the diffusive oxygen supply of the skin from the exterior while
     the emulsion-forming components do not impair the stability of the oxygen
     carrier and its diffusion. The use of the preparation as a cosmetic agent
     allows a natural coloration and an addnl. supply of the skin with
     moisture. Thus a base-emulsion contained (weight/weight%): sorbitan
     monostearate 2.00; Macrogol-9-stearate 3.00; glycerol (85%) 5.00;
     triglycerides, medium chain 5.00; citric acid 0.07; potassium sorbate
     0.14; water to 100. A 30 weight/weight% swine Hb solution contained 50 mM
sodium
     bicarbonate and 150 mM sodium chloride; 16 g of the solution were emulsified
     with 84 g of the base emulsion.
TT
     138-81-8
     RL: COS (Cosmetic use); BIOL (Biological study); USES (Uses)
        (cosmetic emulsions containing Hb and myoglobin as oxygen carriers for
        natural regeneration of skin in case of oxygen deficiency)
RN
     138-81-8 CAPLUS
CN
     Propanoic acid, 2,3-bis(phosphonooxy) - (9CI) (CA INDEX NAME)
      OP03H2
HO_2C-CH-CH_2-OPO_3H_2
L4
     ANSWER 5 OF 10 CAPLUS COPYRIGHT 2004 ACS on STN
AN
     2001:935379 CAPLUS
DN
     136:58832
TI
     Improved injectable dispersions of propofol
IN
     Pace, Gary; Vachon, Michael G.; Mishra, Awadhesh K.; Snow, Robert A.
PA
     RTP Pharma Inc., USA
SO
     PCT Int. Appl., 50 pp.
     CODEN: PIXXD2
DT
     Patent
LA
     English
FAN.CNT 1
     PATENT NO.
                      KIND DATE
                                          APPLICATION NO. DATE
                                          -----
PΙ
     WO 2001097779
                      A2
                            20011227
                                          WO 2001-US19009 20010614
     WO 2001097779
                      A3
                            20020919
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             CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM,
            HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS,
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A sterile, injectable homogenized dispersion of micromatrixes or AΒ microdroplets having a mean diameter from about 50 nm to about 1000 nm comprising about 1-7.5 of propofol, about 1-8 of a propofol-soluble diluent, and about 0.67-5 of a surface stabilizing amphiphilic agent suspended in an aqueous medium containing a synergetic quantity of antimicrobial agent and a tonicity modifying amount of a pharmaceutically acceptable water-soluble hydroxyl-group-containing excipient, wherein the ratio of propofol

to diluent is in the range of about 0.25 to about 7.5 while the ratio of propofol to amphiphilic agent is in the range from about 0.4 to about 1.5, and wherein the viscosity of the dispersion is in the range of 1.1 to 8 cps, processes for the formation of the dispersion, and methods of use are disclosed.

ΙT 111616-41-2

> RL: MOA (Modifier or additive use); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(injectable dispersions of propofol)

RN 111616-41-2 CAPLUS

CN4,6,10,12-Tetraoxa-5,11-diphosphapentadecane-1,2,8,14,15-pentol, 5,11-dihydroxy-, 5,11-dioxide, (2R,14R)-rel- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

- 1.4 ANSWER 6 OF 10 CAPLUS COPYRIGHT 2004 ACS on STN
- AN 2000:220734 CAPLUS
- DN 132:256077
- TI Compositions for sustained release of a antimicrobial gas
- IN Wellinghoff, Stephen T.; Barenberg, Sumner A.; Kampa, Joel J.; Barlow, Darren E.
- PA Bernard Technologies, Inc., USA
- SO U.S., 43 pp., Cont.-in-part of U.S. 5,650,446. CODEN: USXXAM
- DT Patent
- LA English

FAN.	CNT 12					
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE	
ΡI	US 6046243	A	20000404	US 1997-858860	19970519	
	US 5360609	Α	19941101	US 1993-17657	19930212	
	US 5631300	Α	19970520	US 1995-462164	19950605	
	US 5650446	A	19970722	US 1995-465358	19950605	
	US 5668185	A	19970916	US 1995-461716	19950605	

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US 5705092
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     US 5980826
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     WO 9852412
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                                                               19980424
             JP, SG
         W:
         RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL,
             PT, SE
     EP 982986
                        A1
                             20000308
                                             EP 1998-918754
                                                               19980424
         R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, NL, SE, PT, IE, FI
                                                               19980424
                                             JP 1998-550371
     JP 2002507195
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                             20020305
                             19981119
                                             AU 1998-63637
                                                               19980428
     AU 9863637
                        A1
     AU 717604
                             20000330
                        B2
PRAI US 1993-17657
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                             19930212
                        B2
                             19940203
     US 1994-192498
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     US 1994-192499
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     US 1994-228671
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     US 1995-461304
                        A2
                             19950605
     US 1995-461706
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     US 1995-461716
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     US 1993-16904
                       B3
                             19930212
     US 1997-858860
                             19970519
                       Α
     WO 1998-US8387
                             19980424
                       W
OS
     MARPAT 132:256077
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AB A composite for retarding microbiol. contamination containing a hydrophobic material containing an acid releasing agent, and a hydrophilic material containing

anions that are capable of reacting with hydronium ions to generate a gas. The hydrophilic and hydrophobic materials are adjacent and substantially free of water, and the hydrophilic material is capable of generating and releasing the gas after hydrolysis of the acid releasing agent. A composition was prepared containing sodium chlorite, formamide, acrylamide, isopropylacrylamide and hydrophobic material consisting of a 40% solution of maleic anhydride-styrene copolymer in ethylbenzene plasticizer. Hydronium ions formed during hydrolysis reacted with chlorite anions to release chlorine dioxide.

IT 196805-61-5P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(compns. for sustained release of an antimicrobial gas)

RN 196805-61-5 CAPLUS

CN Octadecanoic acid, 4-hydroxy-4-oxido-6-[[[2-oxido-4-[[(1-oxooctadecyl)oxy]methyl]-1,3,2-dioxaphospholan-2-yl]oxy]methyl]-9-oxo-2-[(phosphonooxy)methyl]-3,5,8-trioxa-4-phosphahexacos-1-yl ester (9CI) (CA INDEX NAME)

PAGE 1-B

- (CH₂)₁₆- Me

RE.CNT 40 THERE ARE 40 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 7 OF 10 CAPLUS COPYRIGHT 2004 ACS on STN

AN 1995:928112 CAPLUS

DN 123:340761

TI Preparation of copper, tin, and zinc salts of saccharide derivatives for personal care products.

IN Traudt, Michael David; Waterfield, Philip Christopher

PA Unilever PLC, UK

SO Eur. Pat. Appl., 8 pp. CODEN: EPXXDW

DT Patent

LA English

FAN.CNT 1

PΙ

PATENT NO. KIND DATE APPLICATION NO. DATE

EP 658565 A1 19950621 EP 1993-310259 19931217

R: DE, FR, GB, IT

PRAI EP 1993-310259 19931217

AB Copper, tin, and zinc salts of polyhydroxy compds. having at ≥4 C atoms and ≥1 acid, ester-linked salt-forming substituent, excluding zinc hexosephosphates and stannous glucose-1-phosphate, are claimed. These salts have anti-bacterial activity and are useful for inclusion in personal care compns., particularly in oral care compns., to impart anti-plaque, anti-caries, anti-gingivitis properties thereto. Thus, disodium glucose-6-phosphate and SnCl2 were stirred 30 min. in H2O; MeOH was added to precipitate stannous glucose-6-phosphate. The latter at 6000 ppm gave 88% kill of plaque bacteria.

IT 488-69-7DP, Fructose-1,6-diphosphate, Sn and Cu salts RL: SPN (Synthetic preparation); PREP (Preparation)

(preparation of copper, tin, and zinc salts of saccharide derivs. for personal care products)

RN 488-69-7 CAPLUS

CN D-Fructose, 1,6-bis(dihydrogen phosphate) (9CI) (CA INDEX NAME)

Absolute stereochemistry.

L4 ANSWER 8 OF 10 CAPLUS COPYRIGHT 2004 ACS on STN

AN 1983:467310 CAPLUS

DN 99:67310

TI Phosphorus-31 and carbon-13 nuclear magnetic resonance studies of anaerobic glucose metabolism and lactate transport in Staphylococcus aureus cells

AU Ezra, Fouad S.; Lucas, Donald S.; Mustacich, Robert V.; Russell, Anne F.

CS Miami Valley Lab., Procter and Gamble Co., Cincinnati, OH, 45247, USA

SO Biochemistry (1983), 22(16), 3841-9 CODEN: BICHAW; ISSN: 0006-2960

DT Journal

LA English

High-resolution Fourier transform 31P and 13C NMR were used to probe several AB aspects of glucose metabolism and lactate transport in the gram-pos. bacterium Staphylococcus aureus. The 31P NMR spectra show resonances due to intracellular (Piin) and extracellular orthophosphate (Piex), sugar phosphate, and nucleoside di- and triphosphates. A peak due to teichoic acid was also identified. Its appearance indicates a relatively high degree of mobility in the backbone of this cell wall polymer. The intracellular pH is estimated from the chemical shift of the Piin resonance and is dependent upon the pH of the external medium. A prominent feature of the 31P NMR spectra is the progressive broadening and downfield shift of the Piin resonance that occur when the cells are maintained in an anaerobic environment. Oxygenation causes a narrowing and an upfield shift of the Piin resonance and reverses the trends observed under anaerobic conditions. These line width and chemical shift variations are attributed mainly to a binding of the orthophosphate to paramagnetic ions accumulated by the cells during growth. The ESR spectrum of a perchloric acid extract shows a sextet characteristic of Mn(II) hexaaquo ions. Apparently, the Mn is involved in O2 metabolism 13C NMR spectra obtained from S. aureus cells incubated anaerobically with [1-13C] - or [6-13C] glucose show resonances due to fructose 1,6-diphosphate as an intermediary metabolite and mannitol, lactate, and EtOH as the major end products of glucose metabolism The identity of mannitol is determined from the 13C NMR spectrum of a perchloric acid extract The pH of the external medium affects the glycolytic rate and the distribution of end products. When the pH of the medium is raised from 6.0 to 7.5, the rate of glucose consumption is enhanced, whereas the amount of mannitol produced relative to lactate is drastically The latter effect is explained in terms of the regulation of phosphofructokinase activity by the intracellular pH. The intra- and extracellular lactate appear as 2 well-resolved resonances due primarily to the presence of the Mn2+ inside the cells. The result is a downfield shift and broadening of the intracellular resonance which depend on the oxygenation state of the cells and resemble the trends observed in the 31P NMR spectra. The chemical shift inequivalence of the 2 lactate resonances allows the distribution and transport of this metabolite to be measured, with both the internal and external components being monitored independently. During anaerobic glycolysis, a lactate concentration gradient favoring the cytoplasmic compartment is established. The final intracellular concentration is estimated to be 2-5-fold greater than that in

the

external medium. In the presence of O2, lactate is transported into the cells. A rapid efflux occurs as the cells revert to an anaerobic state. Treatment with a fatty acid antimicrobial agent, octanoate, results in a concentration-dependent reduction of the transmembrane pH gradient and a

loss of lactate from the cells during glycolysis. In addition, the uptake of lactate during oxygenation is completely inhibited.

IT 488-69-7

RL: PROC (Process)

(of Staphylococcus aureus, NMR of)

RN 488-69-7 CAPLUS

CN D-Fructose, 1,6-bis(dihydrogen phosphate) (9CI) (CA INDEX NAME)

Absolute stereochemistry.

L4 ANSWER 9 OF 10 CAPLUS COPYRIGHT 2004 ACS on STN

AN 1983:400197 CAPLUS

DN 99:197

TI Experimental candidiasis in rabbits: protective action of fructose-1,6-diphosphate

AU Tarsi, R.; Simonetti, N.; Orpianesi, C.

CS Inst. Microbiol., Univ. Camerino, Camerino, Italy

SO Mycopathologia (1983), 81(2), 111-16 CODEN: MYCPAH; ISSN: 0369-299X

DT Journal

LA English

GI

AB Fructose-1,6-diphosphate (FDP)(I) [488-69-7] exerts a significant protective action towards Candida albicans infections in rabbits. Such protective action seems related to phagocytic activity stimulation by increased ATP [56-65-5] production

IT 488-69-7

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study)

(antimicrobial activity of, in Candida albicans infection, phagocytosis stimulation and ATP in relation to)

RN 488-69-7 CAPLUS

CN D-Fructose, 1,6-bis(dihydrogen phosphate) (9CI) (CA INDEX NAME)

Absolute stereochemistry.

L4 ANSWER 10 OF 10 CAPLUS COPYRIGHT 2004 ACS on STN

AN 1966:5707 CAPLUS

DN 64:5707

OREF 64:1053e-f

TI Antimicrobial action of sulfurous acid. V. The action of sulfurous acid on the metabolism of respiring and fermenting yeast and Escherichia coli cells

AU Wallnoefer, P.; Rehm, H. J.

CS Deut. Forschungsanstalt Lebensmittelchem., Munich, Germany

SO Zeitschrift fuer Lebensmittel-Untersuchung und -Forschung (1965), 127(4), 195-206

CODEN: ZLUFAR; ISSN: 0044-3026

DT Journal

LA German

cf. CA 63, 10349a. Studies were made on the effects of H2SO3 on the respiration and fermentation metabolism of E. coli and Saccharomyces cerevisiae. H2SO3 inhibits fermentation of S. cerevisiae by blocking NAD-dependent reaction of 3-phosphoglyceraldehyde to 1,3-diphosphoglycerate to a greater degree than that of E. coli. Alc. formation by S. cerevisiae and lactic acid formation by E. coli are not directly inhibited by H2SO3. H2SO3 inhibits respiration of E. coli principally by blocking the NAD-dependent reaction of malate to oxalacetate; through the blocking of the NADP-dependent reaction of isocitrate to oxalacetate; and the NAD-dependent reaction of α -ketoglutarate to S-succinyl-CoA. In vitro, H2SO3 did not inhibit the alc. dehydrogenase of yeast.

IT 1981-49-3, Glyceric acid, anhydride with H3PO4, 3-phosphate (formation from 3-phosphoglyceraldehyde by Saccharomyces cerevisiae, sulfurous acid effect on)

RN 1981-49-3 CAPLUS

CN Propanoic acid, 2-hydroxy-3-(phosphonooxy)-, 1-monoanhydride with phosphoric acid (9CI) (CA INDEX NAME)

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L1

L4

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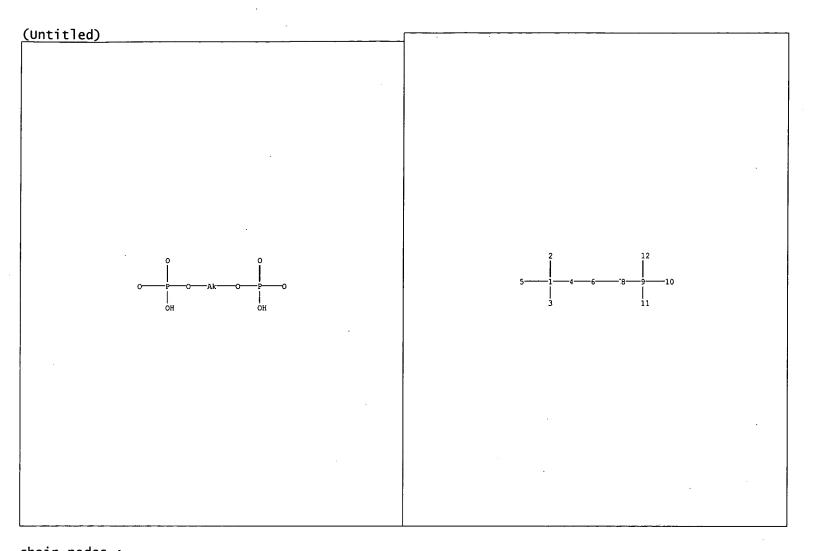
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L3 1425 S L1 SSS FULL

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chain bonds :
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exact bonds :
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normalized bonds :
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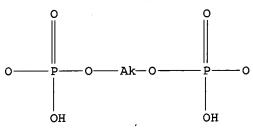
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L1STRUCTURE UPLOADED

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L1 HAS NO ANSWERS

STR L1



Structure attributes must be viewed using STN Express query preparation.

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30.5% PROCESSED

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INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS:

ONLINE **COMPLETE**

COMPLETE BATCH

PROJECTED ITERATIONS:

62127 TO 68993 1796

PROJECTED ANSWERS:

826 TO

L2

20 SEA SSS SAM L1

=> d scan

L220 ANSWERS REGISTRY COPYRIGHT 2004 ACS on STN

Oxirane, (chloromethyl)-, polymer with [(1,1-dimethylethoxy)methyl]oxirane IN and methyloxirane, 2,3-dibromo-1,4-butanediyl bis(dihydrogen phosphate) (9CI)

20 ANSWERS

(C7 H14 O2 . C3 H6 O . C3 H5 Cl O)x . x C4 H10 Br2 O8 P2 MF

CM 1

CM 2

> CM 3

CM 5

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):1

L2 20 ANSWERS REGISTRY COPYRIGHT 2004 ACS on STN

IN D-erythro-L-galacto-2-Nonulose, 1,9-bis(dihydrogen phosphate) (9CI)

MF C9 H20 O15 P2

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):1

L2 20 ANSWERS REGISTRY COPYRIGHT 2004 ACS on STN

IN Diphosphoric acid, 2-(acetyloxy)-1,3-propanediyl ester, ion(2-) (9CI)

MF C5 H12 O16 P4

CI COM

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):1

L2 20 ANSWERS REGISTRY COPYRIGHT 2004 ACS on STN

IN Tetracosanamide, N-[2,3-bis(phosphonooxy)-1-[(phosphonooxy)methyl]eicosyl]-

2-(phosphonooxy)- (9CI)

MF C45 H95 N O17 P4

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HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):1

PAGE 1-A

CM 1

Absolute stereochemistry.

CM 2

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):end

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